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(56) Documents cited

GB 1387954  
GB 1075679  
GB 1019963  
GB 0925927

GB 0516834  
US 4342395  
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(58) Field of search

A5T  
B8C  
B8P  
Selected US specifications from IPC sub-class B65D

## (54) Pack for medicament

(57) A pack (7, 36) provided comprising a circular carrier disc provided with a plurality of containers (8, 37) arranged in a circle and each containing a dose of medicament in particulate form, the containers being openable to permit the medicament therein to be released.

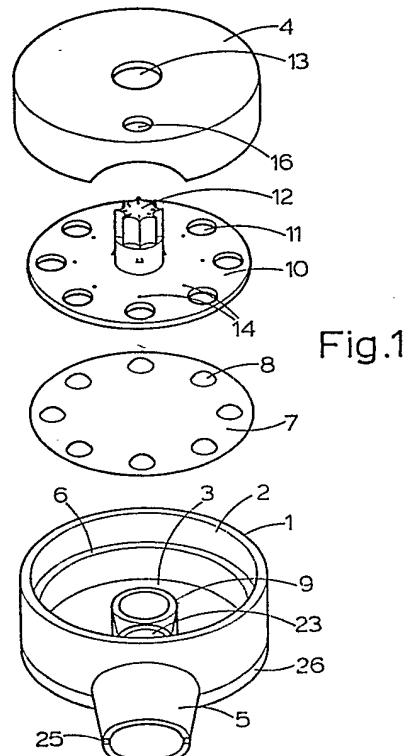


Fig.1

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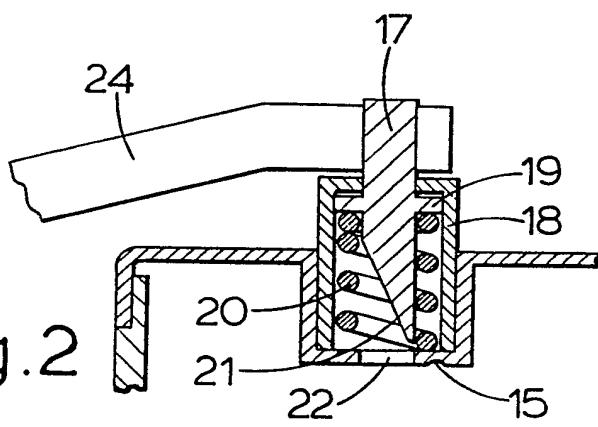
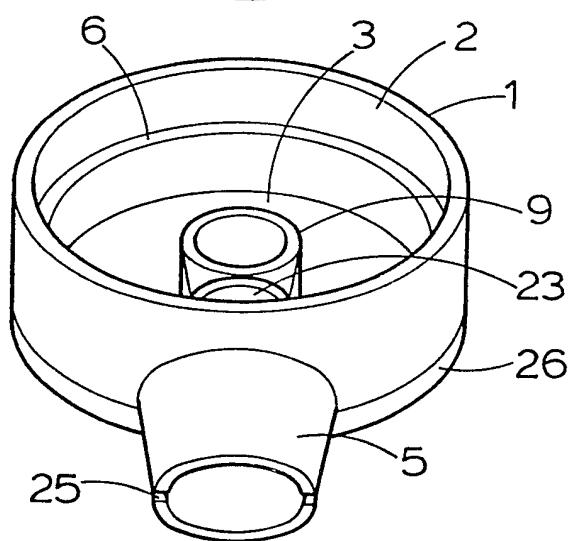
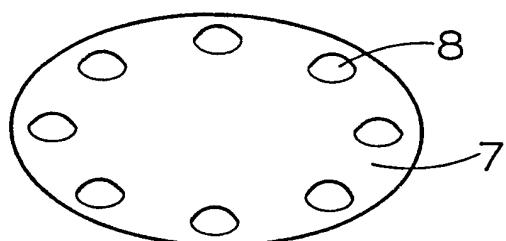
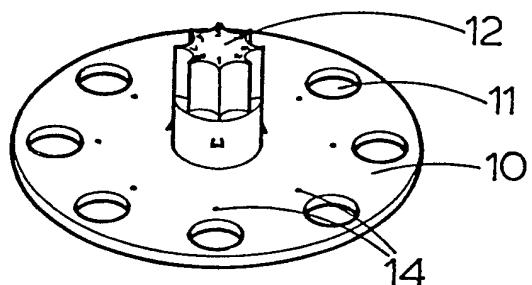
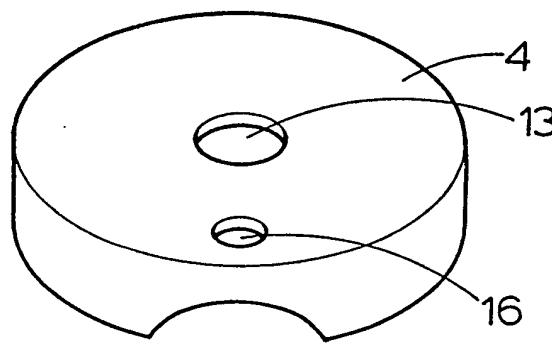


Fig.2

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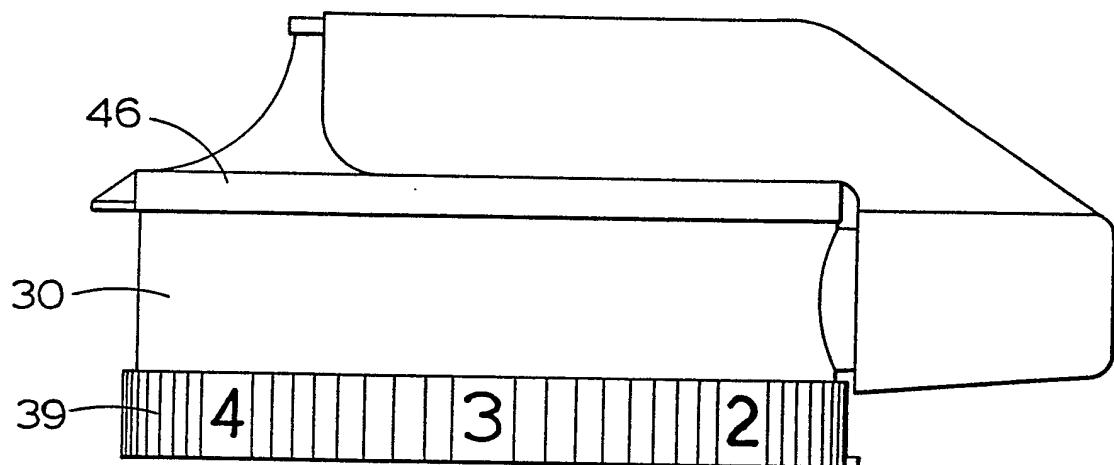


Fig.3

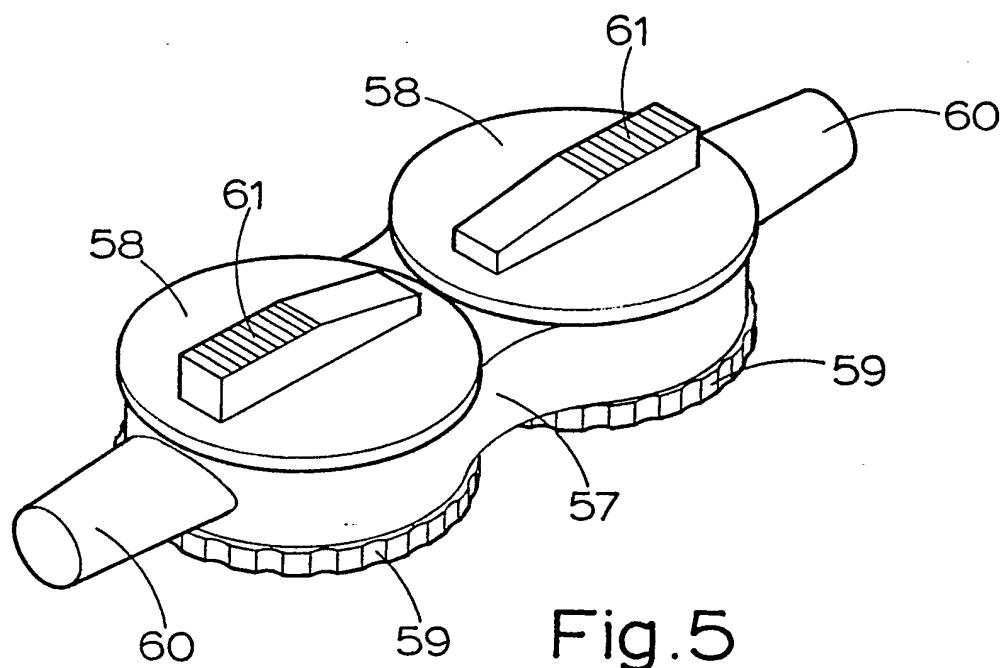


Fig.5

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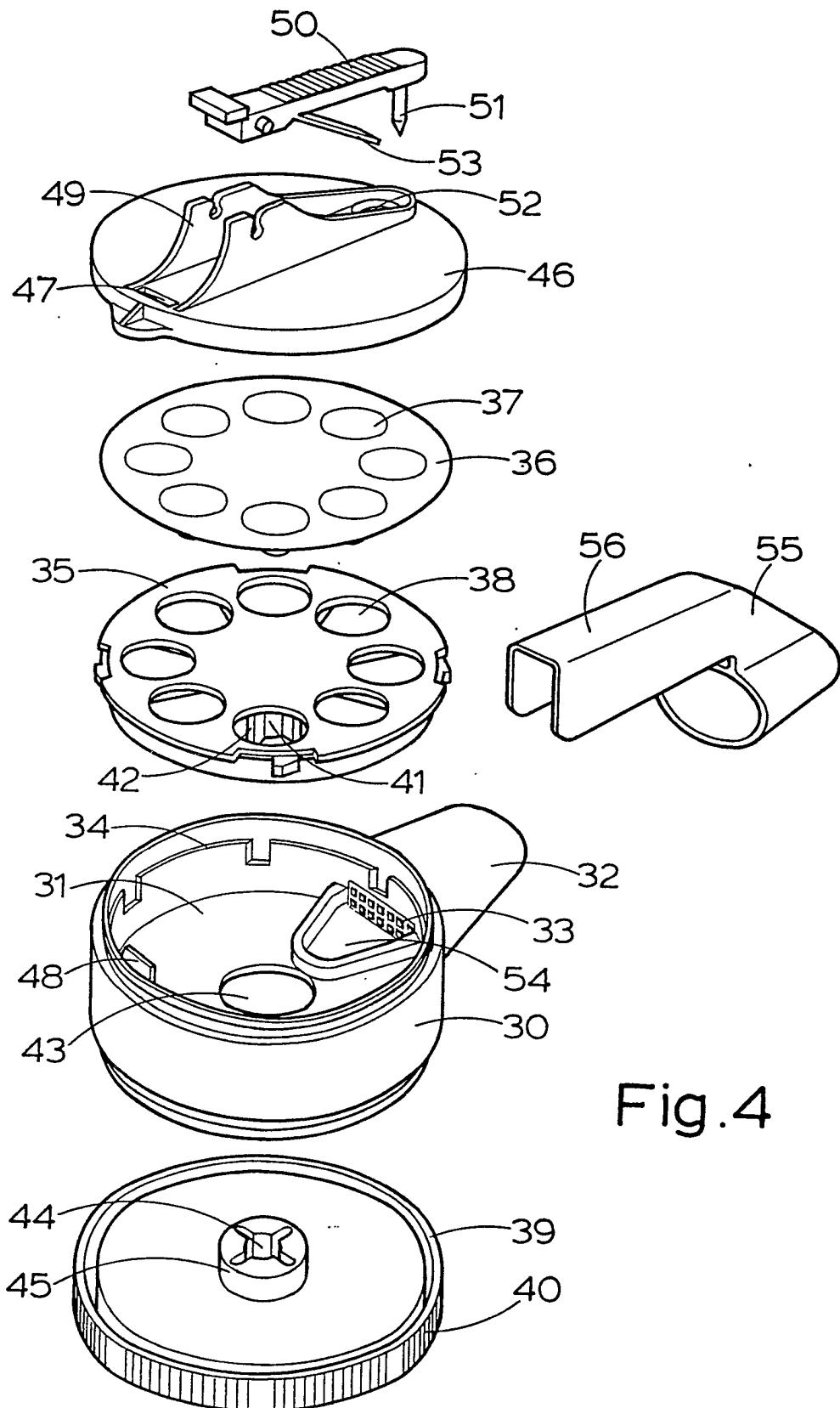


Fig. 4

## SPECIFICATION

### Devices for administering medicaments to patients

5 This invention relates to devices by which a medicament can be administered to or by patients inhaling through the devices. The medicaments may be in solid finely divided form or fluid form. Such devices are now quite well known for administering medicaments contained in capsules to patients suffering from bronchial conditions such as, for example, bronchial asthma. It is well known for medicament in powder or other finely divided form to be supplied in capsules which are loaded by a patient into such a 15 device which is sometimes called an "insufflator". The medicament is then released from the capsule and inhaled by the patient, usually through the mouth, but sometimes through the nose.

The specifications of PCT Application Publication 20 No. WO82/01470 and U.K. Patent Specification No. 1387954 both describe devices for dispensing medicament in finely divided form from capsules. In each of these previously described devices, the capsules are mounted on a rotatable support member on 25 which each capsule in turn can be brought to a position in which it is opened to enable medicament to exit from the capsule to permit it to be inhaled by a patient inhaling through a mouthpiece of the device. In the device described in U.K. Specification 30 No. 1387954, the capsules may be mounted in a so-called blister pack.

There are disadvantages in the use of capsules, which are made of gelatin, to contain medicaments. Gelatin is relatively unstable and is lacking in 35 physical strength so that the capsules need to be protected by packaging, for example in glass bottles. Environmental degradation of both the capsules and their contents may occur in a relatively short time.

An object of the present invention is to provide a 40 more convenient way of administering medicament to such patients than has been possible hitherto and which avoids the need to pack medicaments in capsules. The device of the present invention makes use of the technique of packing medicaments by 45 loading them in blister packs, that is to say packs comprising a sheet, which may be laminated, of foil or plastics material which acts as a carrier and which is provided with a number of breakable or openable containers called "blisters" incorporating a sheet 50 secured on a first sheet to form a cover or lid. Such blister packs are in common use with tablets of one kind or another, but we have discovered that they can also be used with medicaments in finely divided solid form or in liquid form.

55 According to the present invention a device for administering medicaments to patients comprises a housing with a cylindrical chamber therein; an air inlet into the chamber; a support inside the chamber arranged to support a carrier provided with a 60 container for medicament or a plurality of containers arranged in a circle; a plunger operable to engage a container registered therewith to open the container in such a way that air being inhaled by a patient will cause the medicament to be released therefrom; 65 means for rotating a carrier on the support to

register the container, or each of them in turn, with the plunger; and communicating with the interior of the chamber, an outlet through which a patient can inhale whereby medicament will be released from a

70 container and entrained in the air flow produced by the patient so as to pass through the outlet. The outlet is conveniently but not essentially a mouthpiece by which a patient can inhale.

The device of the invention is suitable for administering a variety of medicaments such as, for example, salbutamol, beclomethasone dipropionate and disodium cromoglycate.

Some embodiments of the invention are illustrated in the accompanying schematic drawings in 80 which;

*Figure 1* is an exploded perspective view of a device according to one embodiment of the invention;

*Figure 2* is a detailed view of a plunger device of 85 the same device;

*Figure 3* is an elevation of another embodiment of the invention;

*Figure 4* is an exploded view of the embodiment illustrated in *Figure 3*, and

90 *Figure 5* is a perspective view of yet another embodiment of the invention.

In the embodiment of the invention illustrated in *Figures 1 and 2* of the drawings, a medical administration device comprises a shallow cylindrical housing 1 of a plastics material which has a cylindrical chamber 2 therein. The chamber is closed at one end 3, herein considered the bottom of the chamber, and a removable cover 4 is a close fit over the chamber at the other end.

100 A mouthpiece 5 projects outwardly from the cylindrical wall of the housing 1 and communicates with the interior of the chamber 2. A perforated guard, not shown, is provided in the mouthpiece to prevent any solid particles of an undesirably large 105 size being inhaled by a patient inhaling through the mouthpiece.

A rim or shoulder 6 runs round the inside of the chamber 2 to provide an annular support on which a blister pack 7 may be located.

110 The blister pack 7 can conveniently be a foil laminate with a plurality of frangible containers or "blisters" 8 arranged in a circle. The blisters 8 are filled with medicament in particulate form, having a particle size in the range of 0.5 - 10 microns. The

115 medicament may be with a pharmaceutically acceptable carrier such as lactose or starch in particulate form. Alternatively, the medicament may be in liquid form. The blister pack is of circular disc form, and is removably fitted inside the chamber so that it is

120 replaceable when the individual doses of medicament contained in the blisters have been discharged.

The chamber 2 contains a central open cylindrical support column 9 upstanding from the bottom wall 3 of the chamber. A clamp disc member 10 is removably fitted inside the chamber 2 and has on its

125 underside a plurality of locating pegs, not shown, which engage inside the support column. The clamp member 10 is rotatable inside the chamber. In use, the clamp member is placed on top of a blister pack 7 which has already been loaded into the chamber and

is located on the support shoulder 6. The blister pack 7 is preferably a circular disc of foil laminate material with blisters or containers 8. The clamp member 10 has a plurality of apertures 11 which are arranged in 5 a circle and so spaced from each other that each of them will receive one of the blisters 8 of the blister pack 7. A knob 12 is upstanding from the clamp member 10 and when the lid 4 is fitted on the housing 1 the knob 12 will project through an 10 aperture 13 in the top of the lid 4. This knob can be turned by the patient to rotate the clamp member 10 and since the blisters 8 of the blister pack 7 are located in the apertures 11 in the clamp plate 10 rotation of the clamp member will also rotate the 15 blister pack. A plurality of protuberances or pips 14 are provided on the top of the clamp member 10 and engage in a recess 15, Figure 2, on the underside of the cover 4 to make sure that the clamp plate is correctly aligned in position. As will be seen, the 20 knob 12 is fluted to provide openings between the knob and the hole 13 through which air can enter the chamber 2 from the outside.

The cover 4 also has an aperture 16 in which a 25 plunger 17 contained in a plunger housing 18 can be received. The plunger has an annular shoulder 19 and a spring 20 can bear between the shoulder 19 and the bottom of the plunger housing 18 to urge the plunger into an upper or inoperative position. The plunger may be provided with a knife edge 21 or 30 other means to enable the blister to be opened. When the plunger 17 is depressed against the action of the spring 20, the lower edge portion 21 of the plunger 21 will pass through an aperture 22 in the plunger housing to pass through a blister 8 located 35 in register with the plunger. Such engagement will open the blister, and permit the release of medicament therefrom. This action will so open the blister that when a patient inhales air will pass through the blister, the medicament being entrained in the air 40 flow and exiting through the mouthpiece 5 via a transfer cavity 23 inside the chamber in communication with the mouthpiece 5. By rotation of the knob 12 the clamp member 10 and the blister pack 7 can be rotated to bring each blister in turn into location 45 beneath the plunger. The various protuberances or pips 14 will in turn engage in the recess 15 to make sure that the blister pack is correctly registered with the plunger.

It is not essential that the plunger have a knife 50 21 to open the blister. If desired a needle can be used to perforate the blister or the plunger may have a pointed end or even a blunt end or any other convenient opening means may be used.

The mouthpiece cover can have a locking member 55 24 which can be engaged with the plunger when the device is not in use to prevent accidental actuation of the plunger.

In use, the patient needing a dose of medicament 60 may hold the device with the mouthpiece in his mouth. The patient then depresses the plunger to open the blister and give access to the medicament therefrom and inhales through the mouthpiece so that the medicament will be entrained in the air flow and will enter the lungs of the patient. If desired, the 65 mouthpiece can be provided with air inlet apertures

25 to improve the air flow as the patient inhales.

In a modification not illustrated the underside of the blister pack can be supported on another clamp plate instead of the support rim or shoulder 6.

70 The blister pack is conveniently arranged to provide a sufficient number of individual doses for a patient for use during a convenient period such as one day or more. The housing can be modified by providing an additional chamber, not visible, at the 75 bottom, this additional chamber being closed by a removable cover 26. This additional chamber can be used to store replacement blister packs.

The mouthpiece may, if desired, be arranged so that a patient may use it to inhale through the nose.

80 A modified device which does not use the clamp member 10 is illustrated in Figures 3 and 4. The device of this modification comprises a housing 30 having a chamber 31 therein. A mouthpiece 32 projects outwardly from the cylindrical wall of the 85 housing 30 in a generally radial direction and communicates with the interior of the chamber 31. A perforated guard 33 is provided at the entrance to the mouthpiece 32. A rim or shoulder 34 runs round the inside wall of the chamber 31 to provide an 90 annular support for a support member 35 in the form of a circular plate or disc. This support member is arranged to receive a blister pack 36. The blister pack 36 has a plurality of frangible containers 37 arranged in a circular row. These containers are in the form of 95 "blister" of a generally conical form as clearly shown in Figure 4 and contain a medicament as described with reference to Figure 1. The support member 35 has a plurality of holes 38 equal in number to the number of blisters 37 of the blister pack 36. The

100 conical portion of one blister 37 is located in each of the holes 38 when the device is loaded and in use. An external rotatable member 39 with a knurled edge 40 is located in face contact with the bottom of the housing 30. A spindle or the like 41 with radial

105 projections 42 extends centrally from the support member 35 through a hole 43 in the bottom of the housing 30 and into an opening 44 of complementary shape in a spigot 45 of the member 39. The spigot 45 passes through the hole 43 and the spindle 110 41 and 42 engages in the opening 44 so that rotation of the member 39 will cause similar rotation to the support member 35. A removable cover 46 fits on top of the housing 30. An opening 47 is provided in the cover 46 and engages a projection 48 in the

115 housing 30 so as correctly to locate the cover. The cover 46 carries a bracket 49 on which a lever or trigger 50 is pivotally mounted. A plunger 51 is located on the lever or trigger 50 and extends through a hole 52 in the cover. A spring 53 is 120 provided to bear between the trigger or lever 50 and the top of the cover 46 to urge the lever or trigger upwards.

The hole 52 is so positioned that each hole 38 in the support member 35 will register with this hole as 125 the support member 35 is rotated.

When one of the holes 38 is in register with the hole 52 the trigger 50 can be depressed so that its plunger 51, which may be in the form of a needle, will pierce through the blister 37 located in that hole 130 (i.e. pierce the top and the bottom of the blister)

thereby to permit powder to exit from the blister. Some powder will fall into a tray-like compartment 54 inside the chamber 31. When the patient inhales, air passes through the pierced blister so that powder 5 will be entrained in the airflow and will, with powder from the compartment 54, be withdrawn through the guard 33 and the mouthpiece 32. When the device is not in use, the mouthpiece 32 can be enclosed in a mouthpiece cover or sheath 55 which has a channel-like extension 56 which will engage with the bracket 49 to prevent the plunger 51 being depressed to enter through the hole 37.

When the device is in use and the patient inhales through the mouthpiece 32 it is, of course, essential 15 for air to be able to enter the interior of the chamber 31. Any suitable air inlets can be provided. Conveniently, however, air can enter through the hole 52 the plunger or needle 51 being smaller in diameter than the diameter of the hole 52 so that it serves as 20 an air inlet.

Figure 5 illustrates a modified device which can conveniently be used to administer two different medicaments to a patient at separate times. Treatment of certain patients does require that they inhale 25 two different kinds of medicament. In the device illustrated in Figure 5, a common housing 57 contains two chambers equivalent to the chamber 2 of the embodiment illustrated in Figure 1 and 2 or to the chamber 31 of the embodiment illustrated in 30 Figures 3 and 4. These two chambers are enclosed by removable covers 58 and blister packs contained in the chambers can be rotated in the manner previously described by rotation of knurled wheels, knobs or other members 59. Outlet mouthpieces 60 35 project outwardly from the common housing 57, each one of these outlets 60 leading into one of the chambers enclosed by the common housing. Trigger mechanisms 61 are provided to enable the blisters of the blister packs contained in the chambers to be 40 pierced so that the contents thereof can be inhaled by the patient.

#### CLAIMS

- 45 1. A pack comprising a circular carrier disc provided with a plurality of containers arranged in a circle and each containing a dose of medicament in particulate form, the containers being openable to permit the medicament therein to be released.
- 50 2. A pack as claimed in claim 1 wherein the containers are of generally conical form.
3. A pack as claimed in either preceding claim wherein the carrier disc is of foil laminate material.
4. A pack as claimed in any preceding claim 55 wherein the medicaments is salbutamol or beclomethasone dipropionate in particulate form.
5. A pack as claimed in any one of claims 1 to 3 wherein the medicament is disodium cromoglycate in particulate form.
- 60 6. A pack as claimed in claim 4 or 5 in which the particle size of the medicament is substantially in the range 0.5 - 10 microns.
7. A pack as claimed in any preceding claim in which the medicament is admixed with a solid pharmaceutically acceptable carrier.

8. A pack as claimed in claim 7 in which the carrier is lactose.

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